

Process for preparing 6-alkoxy-(6H)-  
dibenzo[c,e][1,2]oxaphosphorins

The present invention relates to the preparation of 6-  
5 alkoxy-(6H)-dibenzo[c,e][1,2]oxaphosphorins.

The only method disclosed by the literature for  
preparing 6-alkoxy-(6H)-dibenzo[c,e][1,2]oxaphosphorins  
is the alcoholysis of 6-chloro-(6H)-  
10 dibenzo[c,e][1,2]oxaphosphorins in the presence of  
stoichiometric amounts of base such as tertiary amines  
or ammonia (EP 0787738 A1, EP 0304782 A2, Phosphorus  
and Sulfur 1987, 31, p. 71).

15 Reactions of 6H-dibenzo[c,e][1,2]oxaphosphorin 6-oxides  
with orthoformic esters have to date selectively  
afforded only 6-(dialkoxymethyl)-  
dibenzo[c,e][1,2]oxaphosphorin 6-oxides (J. praktische  
Chemie 1979, 321, p. 361).

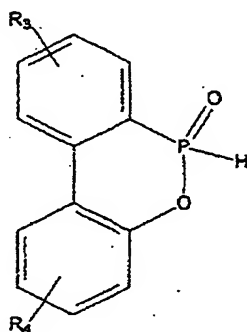
20 The preparation of 6-alkoxy-(6H)-  
dibenzo[c,e][1,2]oxaphosphorins by alcoholysis of 6-  
chloro-(6H)-dibenzo[c,e][1,2]oxaphosphorins by means of  
bases entails a two-stage preparation of the 6-chloro  
25 derivative from o-hydroxybiphenyl and phosphorus  
trichloride with an unsatisfactory overall yield of  
less than 50%. Specifically the second synthetic stage  
is carried out under metal halide catalysis at  
temperatures above 200°C and with HCl elimination. This  
30 process places such high requirements on the technology  
that no industrial solution has to date been attempted.  
In contrast, 6H-dibenzo[c,e][1,2]oxaphosphorin 6-oxide  
is currently even industrially available and is  
prepared from the same reactants in a one-stage  
35 synthesis in yields of above 95% (EP 0806429 A2).

It is thus desirable to develop a process which enables  
the preparation of 6-alkoxy-(6H)-

dibenzo[c,e][1,2]oxaphosphorins by a simple and inexpensive route. They are already known to be useful as additives or modifiers for several polymers and also as intermediates for preparing photoinitiators  
5 (EP-B 0292786, 7856250 and EP-A 0304782).

The object of the present invention is achieved by using 6H-dibenzo[c,e][1,2]oxaphosphorin 6-oxides of the formula I

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as the reactant.

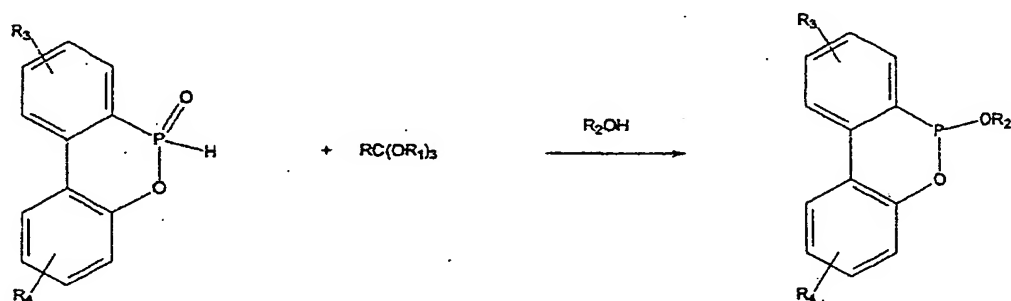
15 In general, the process according to the invention includes the following individual steps: 1) providing at least one solvent, 2) adding the reactant, 3) adding an ortho ester, 4) adding alcohol if it is not already present in the form of the solvent.

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Useful solvents are methanol, ethanol and nonaromatic substituted alcohols, benzene, alkylated benzenes, aliphatic and cycloaliphatic ethers.

25 According to the aforesaid, the present invention relates to a process for gently and selectively preparing 6-alkoxy-(6H)-dibenzo[c,e][1,2]oxaphosphorins by reacting industrially available 6H-dibenzo[c,e][1,2]oxaphosphorin 6-oxides with  
30 orthocarboxylic esters, which is acid-catalyzed. The selection of a suitable alcohol as the reaction medium

then allows the desired 6-alkoxy-(6H)-dibenzo[c,e][1,2]oxaphosphorin to be obtained by in situ transesterification by means of this alcohol. Accordingly, the reaction may be illustrated as follows:



For the  $R_1$  and  $R_2$  radicals in the above formulae, specific substances which may be used include:

optionally substituted alkyl: saturated, straight-chain or branched hydrocarbon radicals, especially having from 1 to 10 carbon atoms, e.g. C1-C6-alkyl such as methyl, ethyl, propyl, 1-methylethyl, butyl, 1-methylpropyl, 2-methylpropyl, 1,1-dimethylethyl, pentyl, 1-methylbutyl, 2-methylbutyl, 3-methylbutyl, 2,2-dimethylpropyl, 1-ethylpropyl, hexyl, 1,1-dimethylpropyl, 1,2-dimethylpropyl, 1-methylpentyl, 2-methylpentyl, 3-methylpentyl, 4-methylpentyl, 1,1-dimethylbutyl, 1,2-dimethylbutyl, 1,3-dimethylbutyl, 2,2-dimethylbutyl, 2,3-dimethylbutyl, 3,3-dimethylbutyl, 1-ethylbutyl, 2-ethylbutyl, 1,1,2-trimethylpropyl, 1,2,2-trimethylpropyl, 1-ethyl-1-methylpropyl and 1-ethyl-2-methylpropyl;

optionally substituted alkenyl: unsaturated, straight-chain or branched hydrocarbon radicals, especially having from 2 to 10 carbon atoms and a double bond in any position, e.g. C2-C6-alkenyl such as ethenyl, 1-propenyl, 2-propenyl, 1-methylethenyl, 1-butenyl, 2-butenyl, 3-butenyl, 1-methyl-1-propenyl, 2-methyl-1-

propenyl, 1-methyl-2-propenyl, 2-methyl-2-propenyl, 1-pentenyl, 2-pentenyl, 3-pentenyl, 4-pentenyl, 1-methyl-1-butenyl, 2-methyl-1-butenyl, 3-methyl-1-butenyl, 1-methyl-2-butenyl, 2-methyl-2-butenyl, 3-methyl-2-butenyl, 1-methyl-3-butenyl, 2-methyl-3-butenyl, 3-methyl-3-butenyl, 1,1-dimethyl-2-propenyl, 1,2-dimethyl-1-propenyl, 1,2-dimethyl-2-propenyl, 1-ethyl-1-propenyl, 1-ethyl-2-propenyl, 1-hexenyl, 2-hexenyl, 3-hexenyl, 4-hexenyl, 5-hexenyl, 1-methyl-1-pentenyl, 2-methyl-1-pentenyl, 3-methyl-1-pentenyl, 4-methyl-1-pentenyl, 1-methyl-2-pentenyl, 2-methyl-2-pentenyl, 3-methyl-2-pentenyl, 4-methyl-2-pentenyl, 1-methyl-3-pentenyl, 2-methyl-3-pentenyl, 3-methyl-3-pentenyl, 4-methyl-3-pentenyl, 1-methyl-4-pentenyl, 2-methyl-4-pentenyl, 3,1,1,2-trimethyl-2-propenyl, 1-ethyl-1-methyl-2-propenyl, 1-ethyl-2-methyl-1-propenyl and 1-ethyl-2-methyl-2-propenyl;

optionally substituted alkynyl: straight-chain or branched hydrocarbon groups, especially having from 2 to 20 carbon atoms and a triple bond in any position, e.g. C2-C6-alkynyl such as ethynyl, 1-propynyl, 2-propynyl, 1-butylnyl, 2-butylnyl, 3-butylnyl, 1-methyl-2-propynyl, 1-pentylnyl, 2-pentylnyl, 3-pentylnyl, 4-pentylnyl, 1-methyl-2-butylnyl, 1-methyl-3-butylnyl, 2-methyl-3-butylnyl, 3-methyl-1-butylnyl, 1,1-dimethyl-2-propynyl, 1-ethyl-2-propynyl, 1-hexynyl, 2-hexynyl, 3-hexynyl, 4-hexynyl, 5-hexynyl, 1-methyl-2-pentylnyl, 1-methyl-3-pentylnyl, 1-methyl-4-pentylnyl, 2-methyl-3-pentylnyl, 2-methyl-4-pentylnyl, 3-methyl-1-pentylnyl, 3-methyl-4-pentylnyl, 4-methyl-1-pentylnyl, 4-methyl-2-pentylnyl, 1,1-dimethyl-2-butylnyl, 1,1-dimethyl-3-butylnyl, 1,2-dimethyl-3-butylnyl, 2,2-dimethyl-3-butylnyl, 3,3-dimethyl-1-butylnyl, 1-ethyl-2-butylnyl, 1-ethyl-3-butylnyl, 2-ethyl-3-butylnyl and 1-ethyl-1-methyl-2-propynyl;

an optionally substituted saturated or mono- or diunsaturated ring which, in addition to carbon atoms, may contain from one to three of the following heteroatoms as ring members: oxygen, sulfur and  
5 nitrogen, for example carbocycles such as cyclopropyl, cyclopentyl, cyclohexyl, cyclopent-2-enyl, cyclohex-2-enyl, 5- to 6-membered, saturated or unsaturated heterocycles containing from one to three nitrogen  
10 atoms and/or one oxygen or sulfur atom, such as 2-tetrahydrofuranyl, 3-tetrahydrofuranyl, 2-tetrahydrothienyl, 3-tetrahydrothienyl, 2-pyrrolidinyl, 3-pyrrolidinyl, 3-isoxazolidinyl, 4-isoxazolidinyl, 5-isoxazolidinyl, 3-isothiazolidinyl, 4-isothiazolidinyl, 5-isothiazolidinyl, 3-pyrazolidinyl, 4-pyrazolidinyl,  
15 5-pyrazolidinyl, 2-oxazolidinyl, 4-oxazolidinyl, 5-oxazolidinyl, 2-thiazolidinyl, 4-thiazolidinyl, 5-thiazolidinyl, 2-imidazolidinyl, 4-imidazolidinyl, 1,2,4-oxadiazolidin-3-yl, 1,2,4-oxadiazolidin-5-yl, 1,2,4-thiadiazolidin-3-yl, 1,2,4-thiadiazolidin-5-yl,  
20 1,2,4-triazolidin-3-yl, 1,3,4-oxadiazolidin-2-yl, 1,3,4-thiadiazolidin-2-yl, 1,3,4-triazolidin-2-yl, 2,3-dihydrofur-2-yl, 2,3-dihydrofur-3-yl, 2,4-dihydrofur-2-yl, 2,4-dihydrofur-3-yl, 2,3-dihydrothien-2-yl, 2,3-dihydrothien-3-yl, 2,4-dihydrothien-2-yl, 2,4-dihydrothien-3-yl, 2,3-pyrrolin-2-yl, 2,3-pyrrolin-3-yl,  
25 2,4-pyrrolin-2-yl, 2,4-pyrrolin-3-yl, 2,3-isoxazolin-3-yl, 3,4-isoxazolin-3-yl, 4,5-isoxazolin-3-yl, 2,3-isoxazolin-4-yl, 3,4-isoxazolin-4-yl, 4,5-isoxazolin-4-yl, 2,3-isoxazolin-5-yl, 3,4-isoxazolin-5-yl, 3,4-isoxazolin-5-yl, 2,3-isothiazolin-3-yl, 3,4-isothiazolin-3-yl, 4,5-isothiazolin-3-yl, 2,3-isothiazolin-4-yl, 3,4-isothiazolin-4-yl, 4,5-isothiazolin-4-yl, 2,3-isothiazolin-5-yl, 3,4-isothiazolin-5-yl, 4,5-isothiazolin-5-yl, 2,3-dihydropyrazol-1-yl, 2,3-dihydropyrazol-2-yl, 2,3-dihydropyrazol-3-yl, 2,3-dihydropyrazol-4-yl, 2,3-dihydropyrazol-5-yl, 3,4-dihydropyrazol-1-yl, 3,4-dihydropyrazol-3-yl, 3,4-dihydropyrazol-4-yl, 3,4-

- 5 dihydropyrazol-5-yl, 4,5-dihydropyrazol-1-yl, 4,5-  
dihydropyrazol-3-yl, 4,5-dihydropyrazol-4-yl, 4,5-  
dihydropyrazol-5-yl, 2,3-dihydrooxazol-2-yl, 2,3-  
dihydrooxazol-3-yl, 2,3-dihydrooxazol-4-yl, 2,3-  
dihydrooxazol-5-yl, 3,4-dihydrooxazol-2-yl, 3,4-  
dihydrooxazol-3-yl, 3,4-dihydrooxazol-4-yl, 3,4-  
dihydrooxazol-5-yl, 3,4-dihydrooxazol-2-yl, 3,4-  
dihydrooxazol-3-yl, 3,4-dihydrooxazol-4-yl, 2-  
piperidinyl, 3-piperidinyl, 4-piperidinyl, 1,3-dioxan-  
10 5-yl, 2-tetrahydropyranyl, 4-tetrahydropyranyl, 2-  
tetrahydrothienyl, 3-tetrahydropyridazinyl, 4-  
tetrahydropyridazinyl, 2-tetrahydropyrimidinyl, 4-  
tetrahydropyrimidinyl, 5-tetrahydropyrimidinyl, 2-  
tetrahydropyrazinyl, 1,3,5-tetrahydrotriazin-2-yl and  
15 1,2,4-tetrahydrotriazin-3-yl, preferably 2-  
tetrahydrofuran-2-yl, 2-tetrahydrothienyl, 2-pyrrolidinyl,  
3-isoxazolidinyl, 3-isothiazolidinyl, 1,3,4-oxazolidin-  
2-yl, 2,3-dihydrothien-2-yl, 4,5-isoxazolin-3-yl, 3-  
piperidinyl, 1,3-dioxan-5-yl, 4-piperidinyl, 2-  
20 tetrahydropyranyl, 4-tetrahydropyranyl;

for the  $R_3$  and  $R_4$  radicals in the above formulae,  
specific substances which may be used include the  
following:

25

alkoxy: straight-chain or branched alkyl groups having  
from one to 30 carbon atoms (as specified above) which  
are bonded to the structure via an oxygen atom (-O-).

30

alkylthio: straight-chain or branched alkyl groups  
having from one to 30 carbon atoms (as specified above)  
which are bonded to the structure via a sulfur atom  
(-S-).

35

Optionally substituted alkyl as described above for  $R_1$   
and  $R_2$ .

Optionally substituted alkenyl as described above for  $R_1$  and  $R_2$ .

Optionally substituted alkynyl as described above for  
5  $R_1$  and  $R_2$ .

An optionally substituted saturated or mono- or diunsaturated ring as described above for  $R_1$  and  $R_2$ .

10 An optionally substituted mono- or bicyclic aromatic ring system which, in addition to carbon atoms may contain from one to four nitrogen atoms or one or two nitrogen atoms and one oxygen or sulfur atom or one oxygen or sulfur atom as ring members, i.e. aryl  
15 radicals such as phenyl and naphthyl, preferably phenyl or 1- or 2-naphthyl, and hetaryl radicals, for example 5-membered heteroaromatic rings containing from one to three nitrogen atoms and/or one oxygen or sulfur atom, such as 2-furyl, 3-furyl, 2-thienyl, 3-thienyl, 1-pyrrolyl, 2-pyrrolyl, 3-pyrrolyl, 3-isoxazolyl, 4-isoxazolyl, 5-isoxazolyl, 3-isothiazolyl, 4-isothiazolyl, 5-isothiazolyl, 1-pyrazolyl, 3-pyrazolyl, 4-pyrazolyl, 5-pyrazolyl, 2-oxazolyl, 4-oxazolyl, 5-oxazolyl, 2-thiazolyl, 4-thiazolyl, 5-thiazolyl, 1-imidazolyl, 2-imidazolyl, 4-imidazolyl, 1,2,4-oxadiazol-3-yl, 1,2,4-oxadiazol-5-yl, 1,2,4-thiadiazol-3-yl, 1,2,4-thiadiazol-5-yl, 1,2,5-triazol-3-yl, 1,2,3-triazol-4-yl, 1,2,3-triazol-5-yl, 1,2,3-triazol-4-yl, 5-tetrazolyl, 1,2,3,4-thiatriazol-5-yl and 1,2,3,4-oxatriazol-5-yl, in particular 3-isoxazolyl, 5-isoxazolyl, 4-oxazolyl, 4-thiazolyl, 1,3,4-oxadiazol-2-yl and 1,3,4-thiadiazol-2-yl;

35 six-membered heteroaromatic rings containing from one to four nitrogen atoms as heteroatoms, such as 2-pyridinyl, 3-pyridinyl, 4-pyridinyl, 3-pyridazinyl, 4-pyridazinyl, 2-pyrimidinyl, 4-pyrimidinyl, 5-pyrimidinyl, 2-pyrazinyl, 1,3,5-triazin-2-yl, 1,2,4-

triazin-3-yl and 1,2,4,5-tetrazin-3-yl, in particular 2-pyridinyl, 3-pyridinyl, 4-pyridinyl, 2-pyrimidinyl, 4-pyrimidinyl, 2-pyrazinyl and 4-pyridazinyl.

- 5 The additional specification "optionally substituted" in relation to alkyl, alkenyl and alkynyl groups is intended to express that these groups may be partially or fully halogenated (i.e. the hydrogen atoms of these groups may be partly or fully replaced by identical or  
10 different halogen atoms as specified above (preferably fluorine, chlorine and bromine, in particular fluorine and chlorine), and/or may bear from one to three, in particular one, of the following radicals:
- 15 nitro, cyano, C1-C4-alkoxy, C1-C4-alkoxycarbonyl or an optionally substituted mono- or bicyclic aromatic ring system which, in addition to carbon atoms may contain from one to four nitrogen atoms or one or two nitrogen atoms and one oxygen or sulfur atom or one oxygen or  
20 sulfur atom as ring members, i.e. aryl radicals such as phenyl and naphthyl, preferably phenyl or 1- or 2-naphthyl, and hetaryl radicals, for example 5-membered heteroaromatic rings containing from one to three nitrogen atoms and/or one oxygen or sulfur atom, such  
25 as 2-furyl, 3-furyl, 2-thienyl, 3-thienyl, 1-pyrrolyl, 2-pyrrolyl, 3-pyrrolyl, 3-isoxazolyl, 4-isoxazolyl, 5-isoxazolyl, 3-isothiazolyl, 4-isothiazolyl, 5-isothiazolyl, 1-pyrazolyl, 3-pyrazolyl, 4-pyrazolyl, 5-pyrazolyl, 2-oxazolyl, 4-oxazolyl, 5-oxazolyl, 2-thiazolyl, 4-thiazolyl, 5-thiazolyl, 1-imidazolyl, 2-imidazolyl, 4-imidazolyl, 1,2,4-oxadiazol-3-yl, 1,2,4-oxadiazol-5-yl, 1,2,4-thiadiazol-3-yl, 1,2,4-thiadiazol-5-yl, 1,2,5-triazol-3-yl, 1,2,3-triazol-4-yl, 1,2,3-triazol-5-yl, 1,2,3-triazol-4-yl, 5-tetrazolyl, 1,2,3,4-thiatriazol-5-yl and 1,2,3,4-oxatriazol-5-yl, in particular 3-isoxazolyl, 5-isoxazolyl, 4-oxazolyl, 4-thiazolyl, 1,3,4-oxadiazol-2-yl and 1,3,4-thiadiazol-2-yl;
- 35



six-membered heteroaromatic rings containing from one to four nitrogen atoms as heteroatoms, such as 2-pyridinyl, 3-pyridinyl, 4-pyridinyl, 3-pyridazinyl, 4-pyridazinyl, 2-pyrimidinyl, 4-pyrimidinyl, 5-pyrimidinyl, 2-pyrazinyl, 1,3,5-triazin-2-yl, 1,2,4-triazin-3-yl and 1,2,4,5-tetrazin-3-yl, in particular 2-pyridinyl, 3-pyridinyl, 4-pyridinyl, 2-pyrimidinyl, 4-pyrimidinyl, 2-pyrazinyl and 4-pyridazinyl.

10

The additional specification "optionally substituted" in relation to the cyclic (saturated, unsaturated or aromatic) groups is intended to express that these groups may be partially or fully halogenated (i.e. the hydrogen atoms of these groups may be partly or fully replaced by identical or different halogen atoms (preferably fluorine, chlorine and bromine, in particular fluorine and chlorine), and/or may bear from one to three of the following radicals: nitro, cyano, C1-C4-alkyl, C1-C4-alkoxy and C1-C4-alkoxycarbonyl.

20

The mono- or bicyclic, aromatic or heteroaromatic systems specified for the radicals may for their part be partially or fully halogenated, i.e. the hydrogen atoms of these groups may be partially or fully replaced by halogen atoms such as fluorine, chlorine, bromine and iodine, preferably fluorine and chlorine.

25

These mono- or bicyclic, aromatic or heteroaromatic systems may, apart from the halogen atoms referred to, additionally bear from one to three of the following substituents:

30

nitro, cyano, thiocyanato;  
alkyl, particularly C1-C6-alkyl as specified above,  
C1-C30-alkoxy,

35

C1-C30-alkylthio,

C1-C4-alkylamino,  
C1-C6-alkylcarbonyl;  
C1-C6-alkoxycarbonyl,  
C1-C6-alkylaminocarbonyl,  
5 C1-C6-alkylcarboxyl,  
C1-C6-alkylcarbonylamino,

C3-C7-cycloalkyl such as cyclopropyl, cyclobutyl,  
cyclopentyl, cyclohexyl and cycloheptyl, preferably  
10 cyclopropyl, cyclopentyl and cyclohexyl, in particular  
cyclopropyl;

C3-C7-cycloalkoxy such as cyclopropyloxy,  
cyclobutyloxy, cyclopentyloxy, cyclohexyloxy and  
15 cycloheptyloxy, preferably cyclopentyloxy and  
cyclohexyloxy, in particular cyclohexyloxy;

C3-C7-cycloalkylthio such as cyclopropylthio,  
cyclobutylthio, cyclopentylthio, cyclohexylthio and  
20 cycloheptylthio, preferably cyclohexylthio,;

C3-C7-cycloalkylamino such as cyclopropylamino,  
cyclobutylamino, cyclopentylamino, cyclohexylamino and  
cycloheptylamino, preferably cyclopropylamino and  
25 cyclohexylamino, in particular cyclopropylamino;

further radicals for optionally substituted mono- or  
bicyclic, aromatic or heteroaromatic radicals:

30 alkenyl, alkynyl, haloalkenyl, haloalkynyl, alkenyloxy,  
alkynyloxy, haloalkenyloxy, haloalkynyloxy, alkenyl-  
thio, alkynylthio, alkylsulfoxy, alkylsulfonyl,  
alkenylsulfoxy, alkynylsulfoxy, alkynylsulfonyl,

35 for the present invention, the solvents used are  
preferably alcohols or alcohol-containing mixtures. The  
alcohol selected is in particular one in which R<sub>1</sub> is  
different from R<sub>2</sub>. Useful solvents also include

benzene, alkylated benzenes, aliphatic and cycloaliphatic ethers.

One advantage in the selection of an alcohol in which  
5  $R_1$  is different from  $R_2$  is that it is possible to determine the target molecule  $R_2$  radical by the alcohol  $R_2$  radical and to use the inexpensive ortho esters.

According to the invention, the reaction is preferably  
10 carried out in the presence of a compound capable of ester formation with the reactant. Useful for this purpose are, for example, ortho esters, in particular trialkyl or triaryl ortho esters or lactone acetals. Preference is accordingly given in accordance with the  
15 invention to trialkyl orthoformates. Very particular preference is given to methyl or ethyl orthoformates.

In the individual steps, catalysts may be added. Useful  
for this purpose are, for example, Lewis acids and  
20 Brønsted acids. Particular mention should be made here of proton donors. Examples are hydrogen halides, phosphoric acids, sulfuric acids, and the like. Preference is given to hydrogen halides, in particular hydrochloric acid. The catalysts are preferably  
25 recycled.

The resulting products are 6-alkoxy- (or 6-aryloxy)-  
(6H)-dibenzo[c,e][1,2]oxaphosphorins. The alkoxy groups  
are preferably methoxy, ethoxy or propoxy radicals.

30 The process according to the invention enables the preparation of 6-alkoxy-(6H)-dibenzo[c,e][1,2]oxaphosphorins, especially when the solvent used is alcohol, directly from industrially available 6H-  
35 dibenzo[c,e][1,2]oxaphosphorin 6-oxides in one synthetic step with excellent yields. It is advantageous that the fine distillation can be

dispensed with under industrial conditions, and a purity of over 96% by GC can be attained in this case.

It is also advantageous that the process according to the invention enables halogen-free working. When, for example, hydrochloric acid is used, it is used merely as a catalyst. In the course of the removal of the excess alcohol, it is also recycled. The consequence thereof is that no halide wastes arise. A further advantage of the present invention is that the reactants used are available inexpensively. Moreover, when the catalysts used are acidic resins, the present invention enables continuous working. Starting from o-hydroxybiphenyl and phosphorus trichloride, it is a two-stage process, whereas three-stage processes are used in the prior art.

The invention is illustrated in detail hereinbelow with reference to the examples:

20

*6-Methoxy-(6H)-dibenzo[c,e][1,2]oxaphosphorin from 6H-dibenzo[c,e][1,2]oxaphosphorin 6-oxide and trimethyl orthoformate in methanol*

25 1.33 mol (287.5 g) of 6H-dibenzo[c,e][1,2]oxaphosphorin 6-oxide and 2.5 ml of conc. HCl are dissolved in 1230 ml of methanol and the mixture is heated to reflux at 85°C (slightly elevated pressure). After 45 min, a further 0.5 ml of conc. HCl is added, and 2.7 mol  
30 (295 ml) of trimethyl orthoformate are subsequently added dropwise within 5 h. During the dropwise addition of the trimethyl orthoformate, in each case 0.5 ml of conc. HCl is added every 30 min. On completion of the reaction, all volatile constituents are removed under  
35 reduced pressure on a rotary evaporator. The yellow, oily residue is distilled in fine vacuum (0.1 mbar). At 130-135°C, the product distills as a colorless, oily

liquid which solidifies slowly after several weeks.  
Yield: 265 g, 87% of theory.

5 *6-Ethoxy-6H-dibenzo[c,e][1,2]oxaphosphorin from 6H-dibenzo[c,e][1,2]oxaphosphorin 6-oxide, ethanol and triethyl orthoformate*

0.2 mol (43.2 g) of 6H-dibenzo[c,e][1,2]oxaphosphorin 6-oxide and 0.5 ml of conc. HCl are dissolved in 352 ml  
10 of ethanol and the mixture is heated to reflux at 90°C (slightly elevated pressure). After 50 min, a further 0.1 ml of conc. HCl is added, and 0.4 mol (59.3 g, 66.5 ml) of triethyl orthoformate is subsequently added dropwise within 4 h. During the dropwise addition of  
15 the triethyl orthoformate, in each case 0.1 ml of conc. HCl is added every 30 min. On completion of the reaction, all volatile constituents are removed under reduced pressure on a rotary evaporator. The yellow, oily residue is distilled in fine vacuum (0.1 mbar). At  
20 135-142°C, the product distills as a colorless, oily liquid (solidified melt  $F_p = 42^\circ\text{C}$ ). Yield: 44.8 g, 92% of theory.

25 *6-Ethoxy-6H-dibenzo[c,e][1,2]oxaphosphorin from 6H-dibenzo[c,e][1,2]oxaphosphorin 6-oxide reactant, ethanol and trimethyl orthoformate*

5.0 mol (1081 g) of 6H-dibenzo[c,e][1,2]oxaphosphorin 6-oxide and 2.0 ml of conc. HCl are dissolved in  
30 4400 ml of ethanol and the mixture is heated to reflux at 95°C (slightly elevated pressure). After 1 h, a further 1.0 ml of conc. HCl is added, and 6.5 mol (689.8 g, 711.1 ml) of trimethyl orthoformate are subsequently added dropwise within 8 h. During the  
35 dropwise addition of the trimethyl orthoformate, in each case 1.0 ml of conc. HCl is added every 30 min. On completion of the reaction, all volatile constituents are removed under reduced pressure on a rotary

evaporator. The yellow, oily residue is distilled in fine vacuum (0.1 mbar). At 135-142°C, the product distills as a colorless, oily liquid (solidified melt  $F_p = 42^\circ\text{C}$ ). Yield: 1001.0 g, 82% of theory.

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*6-Isopropoxy-6H-dibenzo[c,e][1,2]oxaphosphorin from 6H-dibenzo[c,e][1,2]oxaphosphorin 6-oxide reactant, isopropanol and triethyl orthoformate*

- 10 0.28 mol (59.5 g) of 6H-dibenzo[c,e][1,2]oxaphosphorin 6-oxide are dissolved in 600 ml of isopropanol. 0.6 ml of conc. HCl is added and the mixture heated to reflux at 105°C (slightly elevated pressure). After 1 h, a further 0.15 ml of conc. HCl is added, and 0.55 mol
- 15 (81.5 g, 92 ml) of triethyl orthoformate is subsequently added dropwise within 3 h. During the dropwise addition of the triethyl orthoformate, in each case 0.15 ml of conc. HCl is added every 15 min. On completion of the reaction, all volatile constituents
- 20 are removed under reduced pressure on a rotary evaporator. The yellow, oily residue is distilled in fine vacuum (0.1 mbar). At 142-145°C, the product distills as a colorless, oily liquid. Yield: 35.2 g, 78% of theory.